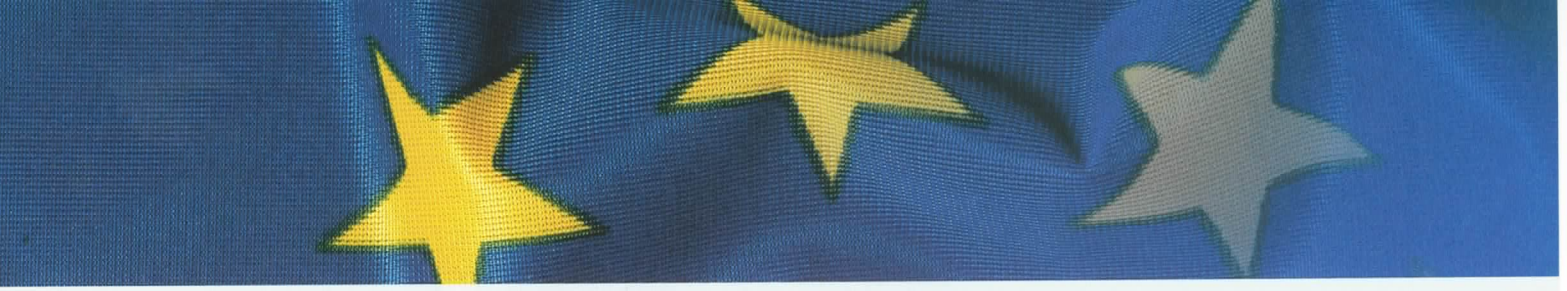

CRDE

CERTIFICATE OF ATTENDANCE

This is to certify that

DANIEL KORCZ

attended and successfully completed the course entitled

AN INTRODUCTION TO CLINICAL TRIALS MONITORING

on 28 – 30 April 2008
in Kraków, Poland

THE SCOPE OF THE COURSE:

Overview of Drug Research and Development
Ethical Issues (Nuremberg Code, Declaration of Helsinki, ICH GCP)
Roles and Responsibilities in Clinical Trial
Informed Consent procedures
Clinical Trial Documentation
Case Report Form
Source Data Verification
Initiation, monitoring, close out visit
Adverse Events, Serious Adverse Events reporting
Workshops: Informed consent review, documentation review,
CRF review, SDV, Initiation Visit, SAE reporting, Monitoring Visit

TRAINING ACCREDITED BY POLISH ASSOCIATION FOR GOOD CLINICAL PRACTICE

Beata Drobniak, Course Leader

30 Apr 2008
Date

Issued by CRDE, Intro/04/08

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